DEPARTMENT OF HEALTH AND HUMAN SERVICES

Public Health Service Food and Drug Administration

APPLICATION FOR A VARIANCE FROM 21 CFR 1040.11(c) FOR A LASER LIGHT SHOW, DISPLAY, OR DEVICE

Form Approved: OMB No. 0910-0025 Expiration Date: October 31, 2000 See Page 4 for OMB Statement.

DOCKET NUMBER

NOTE: No laser light show, projection system, or device may vary from compliance with 21 CFR 1040.11(c) in design or use without the approval of this application in accordance with 21 CFR 1010.4.					
application in accordance with 21 CFR 1010.4. INSTRUCTIONS					
Check all applicable boxes and type or print the	Mail your application to the Dockets Management Branch (HFA-305), Food and				
	orug Administration, Rm 1061, 5630 Fishers Lane, Rockville, MD 20852.				
	nter docket number if assigned.				
1. NAME OF COMPANY Chaotic Laser	Systems				
2. ADDRESS OF COMPANY (Include ZIP Code)(If P.O. Box is used, include	actual street address also.)				
4887 Headlee Dr. Orland	l_0 FC 32822				
3. NAME AND TITLE OF RESPONSIBLE PERSON	4. TELEPHONE NO. (Include area code) 5. DATE OF SUBMISSION				
Joe Settecase Co-Owner	407-277-4611 4/15/99				
THE APPLICANT REQUESTS THE VARIANCE TO BE IN EFFECT FOR	A PERIOD OF (Tro) YEARS FROM THE DATE OF ISSUE. (In				
general, the Agency will approve a variance for only two years. If a longer per	od is requested, a justification must be attached as part of the application.)				
7. PRODUCT DESC	RIPTION AND USE				
a. LIST NAME AND/OR MODEL NUMBER(S) FOR THE LASER LIGHT SHO					
Laser - Lexell 88 Water Cooled Scanner - Redline	lacer's XYP 1000 B "1040.10 and 1040.11)				
b. PRODUCT FOR WHICH A VARIANCE IS REQUESTED	f. PRODUCT IS INTENDED TO BE USED AT ANY ONE LOCATION				
A laser display device	☐ More than 15 days				
A projector for a laser light show	☐ More than 5 but not more than 15 days				
A laser light show	✓ Less than 5 days				
Other (Specify)	g. TOUR IS INTENDED TO RUN FOR				
c. PROJECTORS ARE INTENDED FOR SALE, LEASE, OR LOAN TO	☐ More than 6 months				
OTHER LASER LIGHT SHOW PRODUCERS	☐ 1-6 months				
d. PRODUCT IS INTENDED FOR USE IN A	Less than one month				
☐ Planetarium or other dome projection structure	Not applicable (Not a tour)				
Theater	Other (Specify)				
☐ Hotel/motel ballroom or meeting room	h. PRODUCT UTILIZES THE FOLLOWING LASER EFFECTS				
Store displays	▼ Front screen projections				
☐ Trade show or convention	☐ Rear screen projections				
	☐ Holographic displays				
☐ Discotheque or night club	Multiple reflection/diffraction effects				
Pavilion	Audience scanning (Also includes scanning any accessible)				
Mindoor arena					
▼ Outdoor arena	uncontrolled areas)				
☐ Museum	★ Reflections from stationary mirrors or mirrored surfaces (Beam Matrices)				
Unidoor unenclosed area	Stationary irradiation of rotating mirror balls, etc.				
Other (Specify)	· - /				
e. PRODUCT IS INTENDED TO BE USED	Scanning irradiation of rotating mirror balls, etc.				
☐ At only one (Fixed) location ☐ Fiber optic projections					
At a variety of (Tour) locations **Other (Specify) at a Variety of locations as rental	Fog, smoke, or other scattering enhancement effects				
	ATION LEVELS				
LASER MEDIUM (Ar, He-Ne, etc.) WAVE LE	NGTHS (nm) PEAK POWER (watts)				
1 488 no (13/102)	514 (Green) 5. calts (Runs at 2 myss)				
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O IE ANNUA OCED DADIATION LO DI LOCTO OD COANTED ON ETTE DE C	CHIDATION AND BATE AND SCANNING ERROL ENCY AND AMPLITURE				
9. IF ANY LASER RADIATION IS PULSED OR SCANNED, GIVE THE PULSE DURATION AND RATE AND SCANNING FREQUENCY AND AMPLITUDE					
40. DEACON FOR PEOUFCTING VARIANCE					
10. REASON FOR REQUESTING VARIANCE					
Compliance with the limits of 21 CFR 1040.11(c) would restrict the intended use of the product because compliance would limit the output power to the extent that the desired effects would not be sufficiently visible					
Other or additional explanation (Specify)	1/10/				
1990-1/5/	V/1 K /				
FORM FDA 3147 (7/98) PREVIOUS EDITION IS OBSOLETE Consider to Consider the Consideration of					

11. MANNER IN WHICH IT IS PROPOSED TO DEVIATE FROM THE REQUIREMENTS OF THE APPLICABLE STANDARD It is proposed to deviate from the provisions of 21 CFR 1040.11(c) in that the accessible emission level would exceed the accessible emission limits specified in 21 CFR 1040.11(c).				
	☐ It is proposed to deviate from the provisions of 21 CFR 1040.11(c) as follows:			
12.	ADVANTAGES TO BE DERIVED FROM SUCH DEVIATION			
	Laser light shows and displays are accepted popular media in entertainment and the arts. Use of power levels in excess of the limits imposed by 21 CFR 1040.11(c) is necessary to achieve the required effects in these media.			
	Other or additional advantages (describe and explain).			
13.	EXPLAIN THE ALTERNATE MEANS OF RADIATION PROTECTION TO BE PROVIDED. (Check as many boxes as apply. In item 14 *iustify any boxes not checked, using additional sheets as necessary. State any other means of radiation protection that will be used.)	Remarks,"		
	a. All laser products, systems, shows, and projectors will be certified to comply with 21 CFR 1040.10 and the conditions of this variar be reported as required by 21 CFR 1002.10 AND 1002.11 using the reporting guides provided for such purpose. These actions will accomplished prior to any introduction into commerce.			
	Effects not specifically indicated in this variance application will not be performed. No other effects will be added until an amendmental variance has been obtained and the required reports or supplements, as applicable, have been submitted.	ent to the		
	c. Scanning, projection, or reflection of laser and collateral radiation (Light show radiation) into audience or other accessible uncontrol will not be permitted except for diffuse reflections produced by the atmosphere, added atmospheric scattering media, and target so			
	Laser radiation levels in excess of the limits of Class I will not be permitted at any point less than 3.0 meters above any surface up persons other than operators, performers, or employees are permitted to stand or 2.5 meters below or in lateral separation from an where such persons are permitted to be. Operators, performers, and employees will not be required or allowed to view radiation at limits of Class I or be exposed to radiation above the limits specified in 21 CFR 1040.11(c). We will worth Scan His crowder. Any product which relies on scanning to meet access, exposure, or product class limits will incorporate a scanning safeguard system.	ny place bove the		
	e. Any product which relies on scanning to meet access, exposure, or product class limits will incorporate a scanning safeguard system directly senses scanner motion and which will react fast enough to preclude exceeding the applicable limit.	em which		
	. 📡 All laser light shows shall be under the direct and personal control of trained, competent operator(s). The operator(s) will:			
	(1) Be an employee of the variance holder who will be responsible for the training and the conduct of the operator;			
	(2) Be located where all beam paths can be directly observed at all times; and			
	(3) Immediately terminate the emission of light show radiation in the event of any unsafe condition; or, for outdoor shows, upon reby any air traffic control officials.	quest		
	g. 💢 The maximum laser projector output power will not exceed the level required to obtain the intended effects.			
	The projection system (i.e., the projector and all other components used to produce the lighting effects) will be securely mounted of immobilized to prevent unintended movement or misalignment. Beam masking will be provided as an inherent part of the system of prevent overfilling of screens, beam stops, targets, etc.	r lesign to		
	Laser projectors will not be delivered to any other party under an agreement of sale, lease, or loan unless and until the recipient de that they have a variance in effect at the time of delivery that permits them to produce laser light shows incorporating such projected that they have a variance in effect at the time of delivery that permits them to produce laser light shows incorporating such projected.	emonstrates or(s).		
	In addition to the requirements of 21 CFR 1040.10(h), the manufacturer of laser projectors/systems will provide to parties who pur lease, or borrow the equipment, adequate users' instructions for safe installation and operation which explain the responsibility of as an independent light show manufacturer to submit the required reports and apply for and obtain a variance from CDRH prior to into commerce of any laser light shows.	the recipient		
	The requirements of 21 CFR 1002.30(a)(1) and (2) will be accomplished through the use of written procedures for setup, alignment and performance of each show, these procedures will be in sufficient detail to ensure compliance with 21 CFR 1040.10, the condit variance, and the control of access to radiation areas using the procedures described in the ANSIZ136.1 standard for the safe use (American National Standards Institute, 1430 Broadway, New York, NY 10018) or any other equivalent user consensus standard applicable, state or local requirements. Laser radiation areas which can contain radiation levels above the limits specified in 21 CF will be clearly identified by the posting of warning signs and/or restricting access through physical means (such as pressure switch cells, barriers, guards, etc.). These requirements apply to temporary areas (such as during set up and alignment procedures) and permanent areas. The variance holder will retain the records of these procedures and the results of all tests as required by 21 CFI copy of the variance application, the approval letter, current procedures, and records relating to each particular show will be with the or other responsible individual and will be made available for inspection by FDA and other responsible authorities.	tions of this e of lasers and, where FR 1040.11(c) hes, photo to final or R 1002.31. A		

- I. Advance written notification will be made as early as possible to appropriate federal, state, and local authorities providing show itinerary with dates and locations clearly and completely identified, and a basic description of the proposed effects including a statement of the maximum power output intended. Such notifications will be made, but not necessarily be limited, to:
 - (1) The Center for Devices and Radiological Health, Office of Compliance (HFZ-342), 2098 Gaither Road, Rockville, MD 20850, providing the initial and closing dates for fixed installations and the itinerary for mobile shows. In addition, unless all aspects of each show have been reported and accession numbers clearly referenced, each notice will include detailed descriptions of each show and a listing of all effects to be performed in sufficient detail to confirm compliance with the regulations and this variance.
 - (2) The Federal Aviation Administration (FAA) for any projections into open airspace at any time (i.e., including set up, alignment, rehearsals, performances, etc.). If the FAA objects to any laser effects, the objections will be resolved and any conditions requested by FAA will be adhered to. If these conditions cannot be met, the objectionable effects will be deleted from the show
 - (3) State and local radiation control offices/agencies for all shows to be performed within their jurisdictions. All requirements of state and local law will be satisfied and any objections raised by local authorities will be resolved or the effects deleted. (A list of federal and state offices is available from the Center for Devices and Radiological Health upon request.)

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Our System utilizes a basic Red Line Laser XYP 1000 B beam box which already complies with US CFR 1040.10 and 1040.11. Our shows will consist of basic beam effects to be projected quite a bit higher above the crowd than required. The scanner also contains safely shutter which completely stops the laser beam with the quick press of a button. One of my best friends repairs surgery lasers for a living. In the past year he has taught me a world of information about laser Safety. I also plan to attend a laser safety class part on by Pangolin Laser of Orlando, Ras soon as I my Varience.

CERTIFICATION

I CERTIFY that all of the above information and statements are true, complete, and correct to the best of my knowledge and acknowledge that my variance application may be denied or my variance may be revoked if this application is found to be false, misleading or incorrect in any material way. I have submitted and will submit all reports required by 21 CFR 1002.10 and 1002.11 on the laser equipment and show(s). I further understand that I may be required by regulation or by the Director, Center for Devices and Radiological Health, to supply such other information as may be necessary to evaluate and act on this application.

15. SIGNATURE

16. NAME (Type or Print)

Joe Settecase

17. TITLE

G-Owner

Public reporting burden for this collection of information is estimated to average .5 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to:

DHHS Reports Clearance Officer

Paperwork Reduction Project 0910-0025

Hubert H. Humphrey Building, Room 531-H

200 Independence Avenue, S.W.

Washington, DC 20201

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An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

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www.usps.gov

From: Chaotic Laser Systems 4887 Hendlee Dr. Orlando FC 32822

To: Dockets Management Branch HF-305 Food + Orny Administration Rn 1061 S630 Fishers Lane Rockvilk, MD 20852